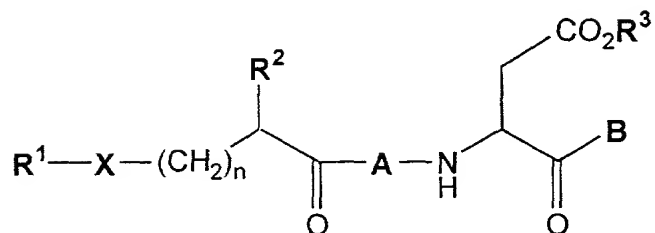


CLAIMS

We claim:

1. A compound of the following formula:



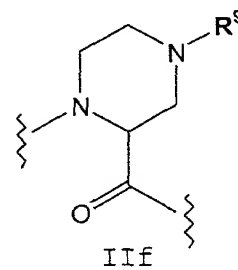
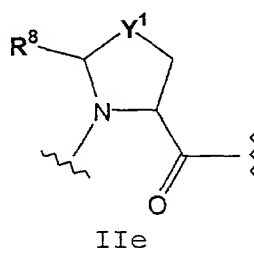
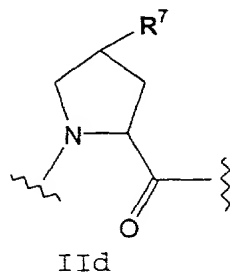
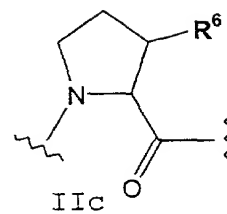
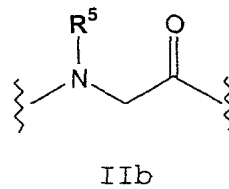
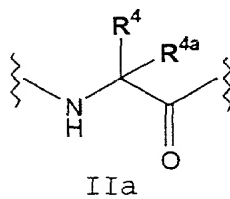
Formula I

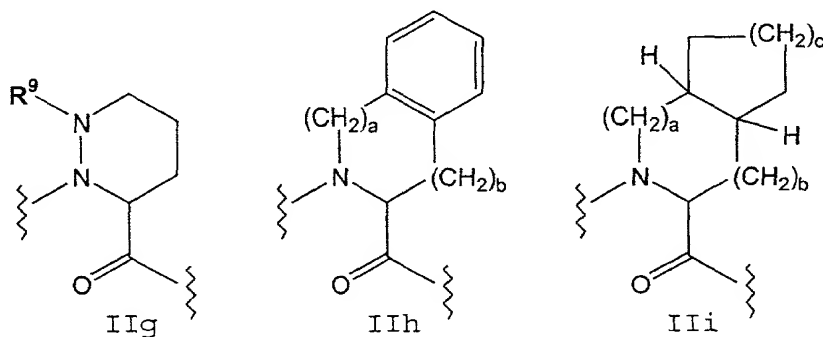
wherein:

n is 0, 1 or 2;

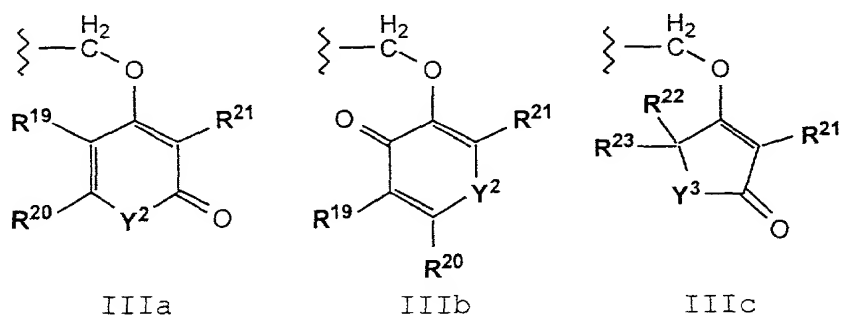
X is CH₂, C=O, O, S or NH;

A is a natural or unnatural amino acid of Formula IIa-i:





B is a hydrogen atom, a deuterium atom, C_{1-10} straight chain or branched alkyl, cycloalkyl, phenyl, substituted phenyl, naphthyl, substituted naphthyl, 2-benzoxazolyl, substituted 2-oxazolyl, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), $(CH_2)_m$ (1 or 2-naphthyl), $(CH_2)_m$ heteroaryl, halomethyl, CO_2R^{13} , $CONR^{14}R^{15}$, CH_2ZR^{16} , $CH_2OCO(aryl)$, $CH_2OCO(heteroaryl)$, or $CH_2OPO(R^{17})R^{18}$, where Z is an oxygen or a sulfur atom, or B is a group of the Formula IIIa-c:



R^1 is phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl, or substituted heteroaryl;

R^2 is hydrogen, alkyl, cycloalkyl, phenyl, substituted phenyl, $(CH_2)_mNH_2$, $(CH_2)_mNHCOR^{10}$, $(CH_2)_mN(C=NH)NH_2$, $(CH_2)_pCO_2R^3$, $(CH_2)_pOR^{11}$, $(CH_2)_pSR^{12}$, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), $(CH_2)_m$ (1 or 2-naphthyl), or $(CH_2)_m$ heteroaryl, wherein heteroaryl includes (but is not limited to)

pyridyl, thienyl, furyl, thiazolyl, imidazolyl, pyrazolyl, isoxazolyl, pyrazinyl, pyrimidyl, triazinyl, tetrazolyl, and indolyl;

R^3 is hydrogen, alkyl, cycloalkyl, (cycloalkyl)alkyl, phenylalkyl, or substituted phenylalkyl;

and wherein

R^4 is alkyl, cycloalkyl, phenyl, substituted phenyl, $(CH_2)_mNH_2$, $(CH_2)_mNHCOR^{10}$, $(CH_2)_mN(C=NH)NH_2$, $(CH_2)_pCO_2R^3$, $(CH_2)_pOR^{11}$, $(CH_2)_pSR^{12}$, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), $(CH_2)_m$ (1 or 2-naphthyl), or $(CH_2)_m$ heteroaryl, wherein heteroaryl includes (but is not limited to) pyridyl, thienyl, furyl, thiazolyl, imidazolyl, pyrazolyl, isoxazolyl, pyrazinyl, pyrimidyl, triazinyl, tetrazolyl, and indolyl;

R^{4a} is hydrogen, or methyl, or R^4 and R^{4a} taken together are $-(CH_2)_d-$ where d is an interger from 2 to 6;

R^5 is phenyl, substituted phenyl, $(CH_2)_p$ phenyl, $(CH_2)_p$ (substituted phenyl), cycloalkyl, or benzofused cycloalkyl;

R^6 is hydrogen, alkyl, cycloalkyl, phenyl, substituted phenyl, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), or $(CH_2)_m$ (1 or 2-naphthyl);

R^7 is hydrogen, fluorine, oxo, alkyl, cycloalkyl, phenyl, substituted phenyl, naphthyl, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), $(CH_2)_m$ (1 or 2-naphthyl), OR^{11} , SR^{12} , or $NHCOR^{10}$;

R^8 is hydrogen, oxo, alkyl, cycloalkyl, phenyl, substituted phenyl, naphthyl, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), or $(CH_2)_m$ (1 or 2-naphthyl);

R^9 is alkyl, cycloalkyl, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), $(CH_2)_m$ (1 or 2-naphthyl), or COR^{10} ;

R^{10} is hydrogen, alkyl, cycloalkyl, phenyl, substituted phenyl, naphthyl, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), $(CH_2)_m$ (1 or 2-naphthyl), OR^{13} , or $NR^{14}R^{15}$;

R^{11} is hydrogen, alkyl, cycloalkyl, phenyl, substituted phenyl, naphthyl, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), or $(CH_2)_m$ (1 or 2-naphthyl);

R^{12} is alkyl, cycloalkyl, phenyl, substituted phenyl, naphthyl, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), or $(CH_2)_m$ (1 or 2-naphthyl);

R^{13} is alkyl, cycloalkyl, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), or $(CH_2)_m$ (1 or 2-naphthyl);

R^{14} is hydrogen, alkyl, cycloalkyl, phenyl, substituted phenyl, naphthyl, substituted naphthyl, $(CH_2)_m$ cycloalkyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), or $(CH_2)_m$ (1 or 2-naphthyl);

R^{15} is hydrogen or alkyl; or

R^{14} and R^{15} taken together form a five, six or seven membered carbocyclic or heterocyclic ring, such as morpholine or N-substituted piperazine;

R^{16} is phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl), $(CH_2)_m$ (1 or 2-naphthyl), or $(CH_2)_m$ heteroaryl;

R^{17} and R^{18} are independently alkyl, cycloalkyl, phenyl, substituted phenyl, naphthyl, or phenylalkyl, substituted phenylalkyl, or (cycloalkyl)alkyl;

R^{19} and R^{20} are independently hydrogen, alkyl, phenyl, substituted phenyl, $(CH_2)_m$ phenyl, or $(CH_2)_m$ (substituted phenyl), or R^{19} and R^{20} taken together are $-(CH=CH)_2-$;

R^{21} is hydrogen, alkyl, phenyl, substituted phenyl, $(CH_2)_m$ phenyl, $(CH_2)_m$ (substituted phenyl);

R^{22} , R^{23} and R^{24} are independently hydrogen or alkyl;

Y^1 is CH_2 , $(CH_2)_2$, $(CH_2)_3$, or S;

Y^2 is O or NR^{24} ;

Y^3 is CH_2 , O, or NR^{24} ;

a is 0 or 1 and b is 1 or 2, provided that when a is 1 then b is 1;

c is 1 or 2, provided that when c is 1 then a is 0 and b is 1;

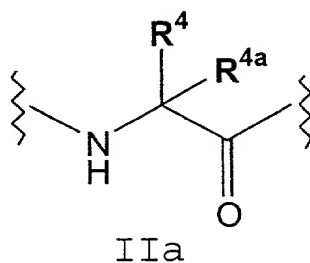
m is 1, 2, 3 or 4; and

p is 1 or 2;

or a pharmaceutically acceptable salt thereof.

2. The compound of claim 1 where X is oxygen.
3. The compound of claim 1 where X is sulfur.
4. The compound of claim 1 where X is NH.
5. The compound of claim 1 where X is CH_2 .
6. The compound of claim 1 where X is C=O.

7. The compound of claim 1 wherein A is

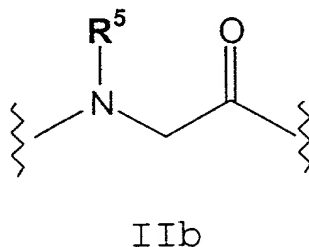


8. The compound of claim 7 wherein

R^4 is lower alkyl, cycloalkyl, phenyl, substituted phenyl, $(CH_2)_nNH_2$, $(CH_2)_mOR^{10}$, $(CH_2)_mSR^{11}$, $(CH_2)_ncycloalkyl$, $(CH_2)_nphenyl$, $(CH_2)_n(substituted\ phenyl)$, or $(CH_2)_n(1\ or\ 2-naphthyl)$; and

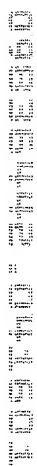
R^{4a} is hydrogen.

9. The compound of claim 1 wherein A is



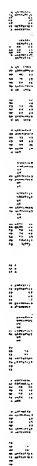
10. The compound of claim 9 wherein R^5 is phenyl, substituted phenyl, $(CH_2)_mphenyl$, $(CH_2)_m(substituted\ phenyl)$, cycloalkyl, or 2-indanyl.

- [illegible]



- [illegible]

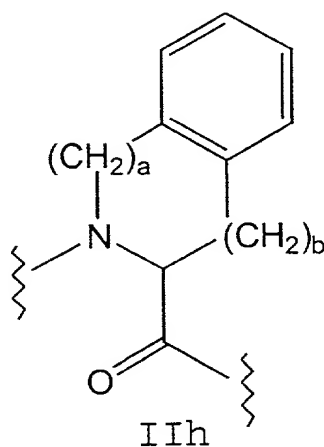
- [illegible]



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[illegible][illegible]

15. The compound of claim 1 wherein A is



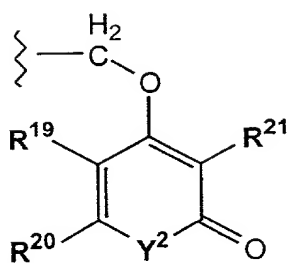
16. The compound of claim 15 wherein a is 0.

17. The compound of claim 1 wherein

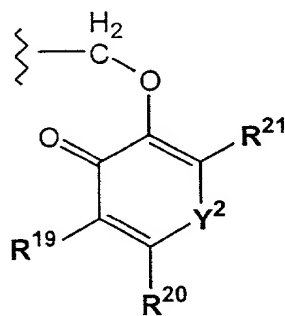
B is hydrogen, 2-benzoxazolyl, substituted 2-oxazolyl, $\text{CH}_2\text{ZR}^{15}$, $\text{CH}_2\text{OCO}(\text{aryl})$, or $\text{CH}_2\text{OPO}(\text{R}^{16})\text{R}^{17}$; and

Z is O or S.

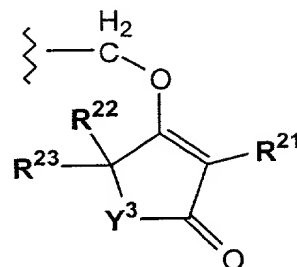
18. The compound of claim 1 wherein B is



IIIa



IIIb



IIIc

19. The compound of claim 18 wherein R^{19} and R^{20} are independently hydrogen, alkyl, or phenyl, or wherein R^{19} and R^{20} taken together are $-(CH=CH)_2-$.

20. The compound of claim 1 wherein

X is O or NH;

n is 0 or 1;

R^1 is substituted phenyl, naphthyl, or substituted naphthyl;

R^2 is hydrogen, lower alkyl, $(CH_2)_pCO_2R^3$, $(CH_2)_m$ (substituted phenyl), $(CH_2)_m$ (1- or 2-naphthyl), or $(CH_2)_m$ tetrazolyl; and

R^3 is hydrogen or lower alkyl.

21. The compound of claim 20 wherein R^1 is 1-naphthyl.

22. The compound of claim 20 wherein R^1 is 2-naphthyl.

23. The compound of claim 20 wherein R^1 is substituted naphthyl.

24. The compound of claim 23 wherein substituted naphthyl is 2-carboxy-1-naphthyl.

25. The compound of claim 20 wherein R^1 is substituted phenyl.

26. The compound of claim 25 wherein substituted phenyl is 2-substituted phenyl.

27. The compound of claim 26 wherein 2-substituted phenyl is (2-phenyl)phenyl.
28. The compound of claim 20 wherein A is alanine, valine, leucine cyclohexylalanine, phenylglycine or t-butylglycine.
29. The compound of claim 28 wherein R^1 is 1-naphthyl.
30. The compound of claim 28 wherein R^1 is 2-naphthyl.
31. The compound of claim 28 wherein R^1 is substituted naphthyl.
32. The compound of claim 31 wherein substituted naphthyl is 2-carboxy-1-naphthyl.
33. The compound of claim 28 wherein R^1 is 2-substituted phenyl.
34. The compound of claim 33 wherein 2-substituted phenyl is (2-phenyl)phenyl.
35. The compound of claim 20 wherein R^2 is $(CH_2)_2CO_2R^3$ and n is 0.
36. A pharmaceutical composition comprising a compound of claim 1 in combination with a pharmaceutically acceptable carrier.

37. A method for treating an autoimmune disease, comprising administering an effective amount of the pharmaceutical composition of claim 36 to a patient in need thereof.

38. A method of treating an inflammatory disease, comprising administering an effective amount of the pharmaceutical composition of claim 36 to a patient in need thereof.

39. A method of treating a neurodegenerative disease, comprising administering an effective amount of the pharmaceutical composition of claim 36 to a patient in need thereof.

40. A method of preventing ischemic injury to a patient suffering from a disease associated with ischemic injury, comprising administering an effective amount of the pharmaceutical composition of claim 36 to a patient in need thereof.

41. A method for expanding of hematopoietic cell populations or enhancing their survival, comprising contacting the cells with an effective amount of the pharmaceutical composition of claim 36.

42. The method of claim 41 wherein the cell populations are granulocytes, monocytes, erythrocytes, lymphocytes or platelets for use in cell transfusions.

43. A method of prolonging the viability of an organ that has been removed from a donor or isolated cells derived from an organ for the purpose of a

[illegible]

- [illegible]